DOCKET NO. 133171.01101 PATENT SERIAL NO. 09/622,452 FILED: October 31, 2000

IN THE CLAIMS:

Please amend claims 1, 7, 12 and 33, cancel claims 4, 9-11, 15, 18, 36, 42, 43, 46, 49, 50, 52-54 and 56-58, and add new claim 59-91 as follows.

This listing of claims will replace all prior versions, and listings of the claims in the application.

Listing of the claims

1. (Currently amended) A pyrogen-free composition comprising a plasmid comprising a nucleotide sequence that encodes an immunogen operably linked to regulatory elements and a nucleotide sequence that encodes an immunomodulating protein operably linked to regulatory elements, wherein said immunomodulating protein is selected from the group consisting of: L-selectin, P-selectin, E-selectin, CD34, GlyCAM-1, MadCAM-1, LFA-1, VLA-1. Mae-1, pl50.95, PECAM, ICAM-2, ICAM-3, CD2, LFA3, mutant forms of IL-18, CD40, CD40L, vascular growth factor, IL-7, nerve growth factor, vascular endothelial growth factor, Fas, TNF receptor, Flt, Apo-1, p55, WSL-1, DR3, TRAMP, Apo-3, AIR, LARD, NGRF, DR4, DR5, KILLER, TRAIL-R2, TRICK2, DR6, and Caspase ICE and wherein said immunogen is a pathogen antigen selected from the group consisting of an influenza antigen, an HIV-1 antigen and an HSV antigen.

2-5. (Canceled)

- 6. (Previously presented) An injectable pharmaceutical composition comprising the pyrogen free composition of claim 1.
- 7. (Currently amended) A method of inducing cytotoxic T cell response in an individual against an immunogen comprising administering by intramuscular injection to said individual a pyrogen free-composition comprising a plasmid comprising a nucleotide sequence that encodes

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an immunogen operably linked to regulatory elements and a nucleotide sequence that encodes an immunomodulating protein operably linked to regulatory elements, wherein said immunomodulating protein is selected from the group consisting of: L. selectin, P. selectin, E-selectin, CD34, GlyCAM-1, MadCAM-1, LFA-1, VLA-1. Mac-1, pl50.95, PECAM, ICAM-2, ICAM-3, CD2, LFA3, mutant forms of IL-18, CD40, CD40L, vascular growth factor, IL-7, nerve growth factor, vascular endothelial growth factor, Fas, TNF receptor, Flt, Apo-1, p55, WSL-1, DR3, TRAMP, Apo-3, AIR, LARD, NGRF, DR4, DR5, KILLER, TRAIL-R2, TRICK2, DR6, and Caspase ICE and wherein said immunogen is a pathogen antigen.

8-11. (Canceled)

12. (Currently amended) A pyrogen-free composition comprising two plasmids: a first plasmid comprising a nucleotide sequence that encodes an immunogen operably linked to regulatory elements; and a second plasmid comprising a nucleotide sequence that encodes an immunomodulating protein operably linked to regulatory elements, wherein said immunomodulating protein is selected from the group consisting of: L. selectin, P. selectin, E. selectin, CD34, GlyCAM-1, MadCAM-1, LFA-1, VLA-1. Mae-1, pl50.95, PECAM, ICAM-2, ICAM-3, CD2, LFA3, mutant forms of IL-18, CD40, CD40L, vascular growth factor, IL-7, nerve growth factor, vascular endothelial growth factor, Fas, TNF receptor, Flt, Apo-1, p55, WSL-1, DR3, TRAMP, Apo-3, AIR, LARD, NGRF, DR4, DR5, KILLER, TRAIL-R2, TRICK2, DR6, and Caspase ICE- and wherein said immunogen is a pathogen antigen selected from the group consisting of an influenza antigen, an HIV-1 antigen and an HSV antigen.

13-16. (Canceled)

17. (Previously presented) An injectable pharmaceutical composition comprising the pyrogen free composition of claim 12.

18-32. (Canceled)

33. (Currently amended) A method of inducing cytotoxic T cell response in an individual against an immunogen comprising administering to said individual by intramuscular injection a composition comprising two plasmids: a first plasmid comprising a nucleotide sequence that encodes said immunogen operable linked to regulatory elements; and a second plasmid nucleic acid molecule comprising a nucleotide sequence that encodes an immunomodulating protein operably linked to regulatory elements, wherein said immunomodulating protein is selected from the group consisting of: L-selectin, P-selectin, E-selectin, CD34, GlyCAM-1, MadCAM-1, LFA-1, VLA-1. Mae-1, pl50.95, PECAM, ICAM-2, ICAM-3, CD2, LFA3, mutant forms of IL-18, CD40, CD40L, vascular growth factor, IL-7, nerve growth factor, vascular endothelial growth factor, Fas, TNF-receptor, Flt, Apo-1, p55, WSL-1, DR3, TRAMP, Apo-3, AIR, LARD, NGRF, DR4, DR5, KILLER, TRAIL-R2, TRICK2, DR6, and Caspase-ICE-and wherein the immunogen is a pathogen antigen.

34-54. (Canceled)

55. (Previously presented) A method of claim 33 wherein said immunogen is a viral antigen.

56-58. (Canceled)

- 59. (New) The pyrogen-free composition of claim 1 wherein said immunogen is an influenza antigen.
- 60. (New) An injectable pharmaceutical composition comprising the composition of claim 59.
- 61. (New) The pyrogen-free composition of claim 1 wherein said immunogen is an HIV-1 antigen.

- 62. (New) An injectable pharmaceutical composition comprising the composition of claim 61.
- (New) The pyrogen-free composition of claim 1 wherein said immunogen is an HSV antigen.
- 64. (New) An injectable pharmaceutical composition comprising the composition of claim 63.
- 65. (New) The method of claim 7 wherein said composition is pyrogen free.
- 66. (New) The method of claim 7 wherein said immunogen is a viral antigen.
- 67. (New) The method of claim 66 wherein said composition is pyrogen free.
- 68. (New) The method of claim 7 wherein the immunogen is selected from the group consisting of: an influenza antigen, an HIV-1 antigen and an HSV antigen.
- 69. (New) The method of claim 68 wherein said composition is pyrogen free.
- 70. (New) The method of claim 7 wherein said immunogen is an influenza antigen.
- 71. (New) The method of claim 70 wherein said composition is pyrogen free.
- 72. (New) The method of claim 7 wherein said immunogen is an HIV-1 antigen.
- 73. (New) The method of claim 72 wherein said composition is pyrogen free.
- 74. (New) The method of claim 7 wherein said immunogen is a HSV antigen.

- 75. (New) The method of claim 74 wherein said composition is pyrogen free.
- 76. (New) The pyrogen-free composition of claim 12 wherein said immunogen is an influenza antigen.
- 77. (New) An injectable pharmaceutical composition comprising the composition of claim 76.
- 78. (New) The pyrogen-free composition of claim 12 wherein said immunogen is an HIV-1 antigen.
- 79. (New) An injectable pharmaceutical composition comprising the composition of claim 78.
- 80. (New) The pyrogen-free composition of claim 12 wherein said immunogen is an HSV antigen.
- 81. (New) An injectable pharmaceutical composition comprising the composition of claim 80.
- 82. (New) The method of claim 33 wherein said composition is pyrogen free.
- 83. (New) The method of claim 55 wherein said composition is pyrogen free.
- 84. (New) The method of claim 33 wherein the immunogen is selected from the group consisting of: an influenza antigen, an HIV-1 antigen and an HSV antigen.
- 85. (New) The method of claim 86 wherein said composition is pyrogen free.
- 86. (New) The method of claim 86 wherein said immunogen is an influenza antigen.

- 87. (New) The method of claim 88 wherein said composition is pyrogen free.
- 88. (New) The method of claim 86 wherein said immunogen is an HIV-1 antigen.
- 89. (New) The method of claim 90 wherein said composition is pyrogen free.
- 90. (New) The method of claim 86 wherein said immunogen is a HSV antigen.
- 91. (New) The method of claim 90 wherein said composition is pyrogen free.